IAP2 Rec'd PCT/PTO 28 SEP 2006

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

APPLICANTS

: MEIKLE, Peter et al.

U.S. Application No.

: Not Yet Assigned

U.S. Application Filed

: Herewith

INT'L APPL'N NO.

: PCT/AU2005/000461

INT'L FILING DATE

: 31 Mar. 2005 (31.03.2005)

FOR

: SCREENING FOR LYSOSOMAL STORAGE DISEASE

STATUS

MS PCT

Commissioner for Patents

P.O. BOX 1450

Alexandria, VA 22313-1450

INFORMATION DISCLOSURE STATEMENT

SIR:

Pursuant to the Duty to Disclose under 37 C.F.R. §1.56(a), applicants enclose herewith a copy of Form PTO-1449 listing documents relevant to the background of the invention described and claimed in the above-identified application. Also enclosed is a copy of the International Search Report in counterpart International Application No. PCT/AU2005/000461.

Applicants respectfully request that the Examiner consider the enclosed references in determining the patentability of the claimed invention. Applicants also request that the Examiner return a copy of enclosed Form PTO-1449 in due course with initials or other markings thereon indicating that the enclosed references have been so considered.

Respectfully submitted,

COLEMAN SUDOL SAPONE, P.C.

Dated: September 27, 2006

Henry D Coleman Reg. Ng. 32,559

714 Colorado Ave.

Bridgeport, Connecticut 06605-1601

203-366-3560

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INFORMATION DISCLOSURE CITATION IN AN APPLICATION Att'y Ref: A20-079 | Serial No: Applicant: MEIKLE, Peter et al. Filing Date: Art Unit:

United States Patent Documents						
Examiner Initial	Document Number	Date	Name	Class	Subclass	Filing Date

Foreign Patent Documents						
Publication Number	Publication Date	Country	Class	Subclass	Translation Yes No	
					1 65	110
		Publication Publication	Publication Publication Country	Publication Publication Country Class	Publication Publication Country Class Subclass	Publication Publication Country Class Subclass Transla

Examiner Initial	Other Documents (by Author, Date, Publication, Pertinent Pages, Title, Etc.)			
	Whitfield PD et al (2002) Mol. Gen & Metabolism 75: 46-55 "Correlation among Genotype, Phenotype and Biochemical Markers in Gaucher Disease: Implications for the Prediction of Disease Severity"			
	Cable WJL et al (1982) Neurology (Ny) 32: 1139-1145 "Fabry disease: Detection of heterozygotes by examination of glycolipids in urinary sediment"			

Examiner:	Date Considered:
EVAMINED: Initial if aitation considered whether or not citat	ion is in conformance with MDED & 600. Draw line through citation if

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Fujiwaki T et al (2002) Brain & Development 24:170-173 "Application of delayed extraction matrix-assisted laser desorption ionization time-of-flight mass spectrometry for analysis of sphingolipids in cultured skin fibroblasts from sphingolipidosis patients"
Fujiwaki T et al (2002) J. Chromatography B 776: 115-123 "Application of delayed extraction-matrix-assisted laser desorption ionization time-of-flight mass spectrometry for analysis of sphingolipids in pericardial fluid, peritoneal fluid and serum from Gaucher disease patients"
Oshima M et al (1990) Bioch et Biophys Ada 1043: 157-160 "Urinary neutral glycosphingolipid analysis of patients with Fabry's disease; rapid isocratic elution from high-performance liquid chromatrography as per-o-benzoyl derivatives"

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